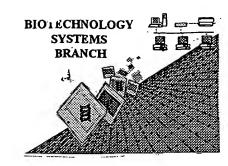
0590

RAW SEQUENCE LISTING ERROR REPORT



The Biotechnology Systems Branch of the Scientific and Technical Information Center (STIC) detected errors when processing the following computer readable form:

| Application Serial Number: | 09/927,8// |
|----------------------------|------------|
| Source: | OIPE |
| Date Processed by STIC: | 10/22/2001 |

THE ATTACHED PRINTOUT EXPLAINS DETECTED ERRORS.
PLEASE FORWARD THIS INFORMATION TO THE APPLICANT BY EITHER:

- 1) INCLUDING A COPY OF THIS PRINTOUT IN YOUR NEXT COMMUNICATION TO THE APPLICANT, WITH A NOTICE TO COMPLY or,
- 2) TELEPHONING APPLICANT AND FAXING A COPY OF THIS PRINTOUT, WITH A NOTICE TO COMPLY

FOR CRF SUBMISSION QUESTIONS, PLEASE CONTACT MARK SPENCER, 703-308-4212.

FOR SEQUENCE RULES INTERPRETATION, PLEASE CONTACT ROBERT WAX, 703-308-4216. PATENTIN 2.1 e-mail help: patin21help@uspto.gov or phone 703-306-4119 (R. Wax) PATENTIN 3.0 e-mail help: patin3help@uspto.gov or phone 703-306-4119 (R. Wax)

TO REDUCE ERRORED SEQUENCE LISTINGS, PLEASE USE THE <u>CHECKER</u> <u>VERSION 3.0 PROGRAM</u>, ACCESSIBLE THROUGH THE U.S. PATENT AND TRADEMARK OFFICE WEBSITE. SEE BELOW:

Checker Version 3.0

The Checker Version 3.0 application is a state-of the-art Windows based software program employing a logical and intuitive user-interface to check whether a sequence listing is in compliance with format and content rules. Checker Version 3.0 works for sequence listings generated for the original version of 37 CFR §§1.821 – 1.825 effective October 1, 1990 (old rules) and the revised version (new rules) effective July 1, 1998 as well as World Intellectual Property Organization (WIPO) Standard ST.25.

Checker Version 3.0 replaces the previous DOS-based version of Checker, and is Y2K-compliant. Checker allows public users to check sequence listings in Computer Readable form (CRF) before submitting them to the United States Patent and Trademark Office (USPTO). Use of Checker prior to filing the sequence listing is expected to result in fewer errored sequence listings, thus saving time and money.

Checker Version 3.0 can be down loaded from the USPTO website at the following address: http://www.uspto.gov/web/offices/pac/checker

OIPE

RAW SEQUENCE LISTING DATE: 10/22/2001 PATENT APPLICATION: US/09/927,811 TIME: 13:03:48

Input Set : A:\ES.txt

19 <170> SOFTWARE: PatentIn Ver. 2.1

Output Set: N:\CRF3\10222001\1927811.raw

ERRORED SEQUENCES

```
494 <210> SEQ ID NO: 27
495 <211> LENGTH: 26
496 <212> TYPE: DNA
497 <213> ORGANISM: Artificial sequence
499 <220> FEATURE:
500 <223> OTHER INFORMATION: Description of the artificial sequence:
sequencing primer Plasm. R (backwards)
503 <400> SEQUENCE: 27
504 ctgctcgctt cgctacttgg agccac

E--> 507
E--> 509
W--> 511
A:\Sequence Listing for Grunecker.doc

A:\Sequence Listing for Grunecker.doc
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sel PiZ

| <210> | 2 |
|--------|--|
| <211> | 15 |
| <212> | DNA |
| <213> | Artificial sequence |
| <220> | |
| <223> | Description of the artificial sequence: |
| | Consensus sequence for a heat schock element |
| <400> | 2 Shock -) global muspelling |
| ngaanr | innnn ngaa(h) |
| · \ | |
| | 1 9 a Sans Ayang Sheet |
| / | sel tim 9 on Ever Summary Sheet |
| | |

Use of n and/or Xaa has been detected in the Sequence Listing. Review the Sequence Listing to insure a corresponding explanation is presented in the <220> to <223> fields of each sequence using n or Xaa.

VERIFICATION SUMMARY

DATE: 10/22/2001 PATENT APPLICATION: US/09/927,811 TIME: .13:03:49

Input Set : A:\ES.txt

Output Set: N:\CRF3\10222001\1927811.raw

L:11 M:270 C: Current Application Number differs, Replaced Application Number L:12 M:271 C: Current Filing Date differs, Replaced Current Filing Date L:52 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:2 L:52 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:2 L:52 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:2 L:65 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:3 L:65 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:3 L:65 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:3 L:288 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:9 L:288 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:9 L:288 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9 L:507 M:254 E: No. of Bases conflict, LENGTH:Input:1 Counted:26 SEQ:27 M:254 Repeated in SeqNo=27 L:511 M:334 W: (2) Invalid Amino Acid in Coding Region, NUMBER OF INVALID KEYS:4 L:512 M:336 W: Invalid Amino Acid Number in Coding Region, SEQ ID:27 L:512 M:334 W: (2) Invalid Amino Acid in Coding Region, NUMBER OF INVALID KEYS:4

Raw Sequence Listing Error Summary

| | 20/2024 |
|-------------------------------------|---|
| ERROR DETECTED | SUGGESTED CORRECTION SERIAL NUMBER: 09/92//8// |
| ATTN: NEW RULES CASI | ES: PLEASE DISREGARD ENGLISH "ALPHA" HEADERS, WHICH WERE INSERTED BY PTO SOFTWAF |
| lWrapped Nucleics Wrapped Aminos | The number/text at the end of each line "wrapped" down to the next line. This may occur if your file was retrieved in a word processor after creating it. Please adjust your right margin to .3; this will prevent "wrapping." |
| 2Invalid Line Length | The rules require that a line not exceed 72 characters in length. This includes white spaces. |
| 3Misaligned Amino Numbering | The numbering under each 5th amino acid is misaligned. Do not use tab codes between numbers; use space characters, instead. |
| 4Non-ASCII | The submitted file was not saved in ASCII(DOS) text, as required by the Sequence Rules. Please ensure your subsequent submission is saved in ASCII text. |
| 5Variable Length | Sequence(s) contain n's or Xaa's representing more than one residue. Per Sequence Rules, each n or Xaa can only represent a single residue. Please present the maximum number of each residue having variable length and indicate in the <220>-<223> section that some may be missing. |
| 6PatentIn 2.0 "bug" | A "bug" in PatentIn version 2.0 has caused fire <220>-<223> section to be missing from amino acid sequences(s) Normally, PatentIn would automatically generate this section from the previously coded nucleic acid sequence. Please manually copy the relevant <220>-<223> section to the subsequent amino acid sequence. This applies to the mandatory <220>-<223> sections for Artificial or Unknown sequences. |
| 7Skipped Sequences (OLD RULES) | Sequence(s) missing. If intentional, please insert the following lines for each skipped sequence: (2) INFORMATION FOR SEQ ID NO:X: (insert SEQ ID NO where "X" is shown) (i) SEQUENCE CHARACTERISTICS: (Do not insert any subheadings under this heading) (xi) SEQUENCE DESCRIPTION:SEQ ID NO:X: (insert SEQ ID NO where "X" is shown) This sequence is intentionally skipped |
| | Please also adjust the "(ii) NUMBER OF SEQUENCES:" response to Include the skipped sequences. |
| 8Skipped Sequences' '(NEW RULES) | Sequence(s) missing. If Intentional, please insert the following lines for each skipped sequence. <210> sequence id number <400> sequence id number 000 |
| 9 Use of n's or Xaa's (NEW RULES) | Use of n's and/or Xaa's have been detected in the Sequence Listing. Per 1.823 of Sequence Rules, use of <220>-<223> is MANDATORY if n's or Xaa's are present. In <220> to <223> section, please explain location of n or Xaa, and which residue n or Xaa represents. |
| 10Invalid <213> Response | Per 1.823 of Sequence Rules, the only valid <213> responses are: Unknown, Artificial Sequence, or scientific name (Genus/species). <220>-<223> section is required when <213> response is Unknown or is Artificial Sequence |
| 11Use of <220> | Sequence(s) missing the <220> "Feature" and associated numeric identifiers and responses. Use of <220> to <223> is MANDATORY if <213> "Organism" response is "Artificial Sequence" or "Unknown." Please explain source of genetic material in <220> to <223> section. (See "Federal Register," 06/01/1998, Vol. 63, No. 104, pp. 29631-32) (Sec. 1.823 of Sequence Rules) |
| Patentin 2.0 "bug" | Please do not use "Copy to Disk" function of PatentIn version 2.0. This causes a corrupted file, resulting in missing mandatory numeric identifiers and responses (as indicated on raw sequence listing). Instead, please use "File Manager" or any other manual means to copy file to floppy disk. |
| 13Misuse of n | n can only be used to represent a single nucleotide in a nucleic acid sequence. N is not used to represent any value not specifically a nucleotide. |

AMC/MH - Biotechnology Systems Branch - 08/21/2001